

Docket No.: 1254-0321PUS1

Confirmation No.: N/A

Examiner: Not Yet Assigned

Art Unit: N/A

(PATENT)

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of: Takaji WAKITA et al.

Application No.: 10/589,902

Filed: August 17, 2006

For: NUCLEIC ACID CONSTRUCT CONTAINING

FULLLENGTH GENOME OF HUMAN HEPATITIS C VIRUS, RECOMBINANT FULLENGTH VIRUS GENOME-REPLICATING CELLS HAVING THE

NUCLEIC ACID CONSTRUCT

TRANSFERRED THEREINTO AND METHOD

OF PRODUCING HEPATITIS C VIRUS

PARTICLE

LETTER

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

Subsequent to the filing of the above-identified application on August 17, 2006, attached hereto is an English Translation of the International Preliminary Report on Patentability issued by the International Bureau on behalf of the International Searching Authority. Please make this document of record for the above-identified application.

Application No.: 10/589,902 Docket No.: 1254-0321PUS1

If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or to credit any overpayment to Deposit Account No. 02-2448 for any additional fees required under 37 C.F.R. § 1.16 or under 37 C.F.R. § 1.17; particularly, extension of time fees.

Dated: December 13, 2006

Respectfully submitted,

Andrew D. Meikle

Registration No.: 32,868

BIRCH, STEWART, KOLASCH & BIRCH, LLP

8110 Gatehouse Road

Suite 100 East P.O. Box 747

Falls Church, Virginia 22040-0747

(703) 205-8000

Attorney for Applicant

Attachment(s)

2 ADM/tmh

PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

NOTIFICATION OF TRANSMITTAL OF COPIES OF TRANSLATION OF THE INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (CHAPTER I OR CHAPTER II OF THE PATENT COOPERATION TREATY)

(PCT Rules 44bis.3(c) and 72.2)

To:

HIRAKI, Yusuke Kamiya-cho MT Bldg. 19F 3-20, Toranomon 4-chome Minato-ku, Tokyo 1050001 **JAPON**



Date of mailing (day/month/year) 28 September 2006 (28.09.2006)	
Applicant's or agent's file reference PH-2372-PCT	IMPORTANT NOTIFICATION
International application No. PCT/JP2005/003232	International filing date (day/month/year) 21 February 2005 (21.02.2005)
Applicant TOKYO METROPOLITAN O	RGANIZATION FOR MEDICAL RESEARCH et al

1.	Transmi	Fransmittal of the translation to the applicant.		
	✓	The International Bureau transmits herewith a copy of the English translation of the international preliminary report or patentability (Chapter I).		
		The International Bureau transmits herewith a copy of the English translation of the international preliminary report of		

patentability (Chapter II).

Transmittal of the copy of the translation to the designated or elected Offices.

The International Bureau notifies the applicant that copies of that translation have been transmitted to the following designated or elected Offices requiring such translation:

3

The following designated or elected Offices, having waived the requirement for such a transmittal at this time, will receive copies of that translation from the International Bureau only upon their request:

AE, AG, AL, AM, AP, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EA, EC, EE, EG, EP, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OA, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

3. Reminder regarding translation into (one of) the official language(s) of the elected Office(s).

The applicant is reminded that, where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary report on patentability (Chapter II).

It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned within the applicable time limit (Rule 74.1). See Volume II of the PCT Applicant's Guide for further details.

> The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland

Authorized officer

Yoshiko Kuwahara

Facsimile No. +41 22 338 82 70

Facsimile No. +41 22 338 82 70

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter I of the Patent Cooperation Treaty)

(PCT Rule 44bis)

Applicant's or agent's file reference PH-2372-PCT	FOR FURTHER ACTION	Sec item 4 below
International application No. PCT/JP2005/003232	International filing date (day/month/year) 21 February 2005 (21.02.2005)	Priority date (day/month/year) 20 February 2004 (20.02.2004)
International Patent Classification (8) See relevant information in Form	h edition unless older edition indicated) PCT/ISA/237	
Applicant TOKYO METROPOLITAN ORGA	NIZATION FOR MEDICAL RESEARCH	

1.	This international preliminary report on patentability (Chapter I) is issued by the International Bureau on behalf of the International Searching Authority under Rule 44 bis.1(a).			
2.	This REPORT consists of a total of 5 sheets, including this cover sheet.			
	In the attached sheets, any reference to the written opinion of the International Searching Authority should be read as a reference to the international preliminary report on patentability (Chapter I) instead.			
3.	3. This report contains indications relating to the following items:			
	Box No. I	Basis of the report		
	Box No. II	Priority		
	Box No. ΠΙ	Non-establishment of opinion with regard to novelty, inventive step and industrial applicability		
	Box No. IV	Lack of unity of invention		
	Box No. V	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement		
	Box No. VI	Certain documents cited		
	Box No. VΠ	Certain defects in the international application		
	Box No. VIII	Certain observations on the international application		
4.	4. The International Bureau will communicate this report to designated Offices in accordance with Rules 44bis.3(c) and 93bis.1 but not, except where the applicant makes an express request under Article 23(2), before the expiration of 30 months from the priority date (Rule 44bis.2).			

Date of issuance of this report 19 September 2006 (19.09.2006)

Yoshiko Kuwahara

Authorized officer

e-mail: pt07@wipo.int

Facsimile No. +41 22 338 82 70 Form PCT/IB/373 (January 2004)

The International Bureau of WIPO 34, chemin des Colombettes

1211 Geneva 20, Switzerland

PATENT COOPERATION TREATY

From the INTERNATIONAL SEARCHING AUTHORI	TY		MNSI
To:			PCT PCT
			ITTEN OPINION OF THE ONAL SEARCHING AUTHORITY
			(PCT Rule 43bis.1)
		Date of mailing (day/month/year)	
Applicant's or agent's file reference		FOR FURTHER A	ACTION
PH-2372-PCT			See paragraph 2 below
International application No. PCT/JP2005/003232	International filing date (e	lay/month/year)	Priority date (day/month/year) 20.02.2004
International Patent Classification (IPC) or both	national classification and	HPC	
Applicant			
TOKYO METROPOLITAN OR	GANIZATION 1	FOR MEDICA	L RESEARCH
This opinion contains indications relations	ing to the following items:	:	
Box No. I Basis of the			
Box No. II Priority	•		·
	shment of opinion with reg	ard to novelty, inventi	ive step and industrial applicability
	y of invention	•	
Reasoned sta		I(a)(i) with regard to a	novelty, inventive step or industrial ement
Box No. VI Certain docu			
Box No. VII Certain defe	cts in the international app	olication	
	ervations on the internation		·
Box No. VIII Calain one	•		
2. FURTHER ACTION If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered.			
If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of 3 months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later.			
For further options, see Form PCT/ISA/220.			
3. For further details, see notes to Form	PCT/ISA/220.		
Name and mailing address of the ISA/JP		Authorized officer	
Name and maining address of the 1989s			
Facsimile No.		Telephone No.	

WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.

PCT/JP2005/003232

Bo	No. 1	Basis of this opinion
J.	With filed.	regard to the language, this opinion has been established on the basis of the international application in the language in which it was unless otherwise indicated under this item.
		This opinion has been established on the basis of a translation from the original language into the following language, which is the language of a translation furnished for the purposes of international search (under
	-	Rule 12.3 and 23.1(b)).
2.	With inver	regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed ation, this opinion has been established on the basis of:
	a.	type of material
ļ		a sequence listing
ļ		table(s) related to the sequence listing
	b.	format of material
l		in written format
		in computer readable form
	¢.	time of filing/furnishing
		contained in the international application as filed.
		filed together with the international application in computer readable form.
		furnished subsequently to this Authority for the purposes of search.
3.	\boxtimes	In addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
4.	Addi	itional comments:
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WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.
PCT/JP2005/003232

Box No. V	Reasoned statemer citations and expla	t under Ru nations sup	de 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; oporting such statement	
1. Statement				
Noveli	y (N)	Claims	1-16, 19-25	YES
		Claims	17, 18, 26	NO
lnvent	ive step (IS)	Claims		YES
		Claims	1-26	NO NO
Indust	rial applicability (IA)	Claims	1-26	YES
		Claims		NO

2. Citations and explanations:

Document 1: Hepatitis C Virus-like Particles Induce Virus-specific Humoral and Cellular Immune Responses in Mice, (M. Lechmann, et al.), Hepatology, 2001, Vol. 34, pages 417-423

Document 2: JP, 2002-171978, A (Tokyo Metropolitan Organization for Medical Research), 18 June, 2002 (18.06.02), full text (Family: none)

Document 3: Inducible System in Human Hepatoma Cell Lines for Hepatitis C Virus Production, (S.P. Lim, et al.), Virology, 2002, Vol. 303, pages 79-99

Document 4: Selectable Subgenomic and Genome-length Dicistronic RNAs Derived from an Infectious Molecular Clone of the HCV-N Strain of Hepatitis C Virus Replicate Efficiently in Cultured Huh7 Cells, (M. Ikeda, et al.), J. Virol., 2002, Vol. 76, pages 2997-3006

The subject matters of claims 17, 18 and 26 do not appear to be novel or to involve an inventive step in view of document 1 cited in the ISR.

Document 1 mentions that HCV-like particles containing the core, E1 and E2 can induce liquid immunity.

Considering that the phrase, "its portion", in claims 17 and 18, and the antibody described in claims 2 and 6, do not clearly specify antigens, the subject matters of the said claims are not distinguishable from the invention described in document 1.

The subject matters of claims 1-26 do not appear to involve an inventive step in view of documents 1-4 cited in the ISR.

It is considered that document 2 describes the genome of hepatitis C virus of genotype 2a. It is considered that document 3 mentions that, by transforming host cells by using replicon RNA where the genome of hepatitis C virus of genotype 1b is under control of a tetracycline-inducible promoter, infectious hepatitis C virus particles and cell strains capable of releasing such virus particles on a stable basis were obtained. It is considered that document 4 mentions that, by introducing a genome of hepatitis C virus at a downstream point in relation to an IRES sequence and a marker gene, the said genome can be replicated in host cells.

In view of the foregoing, a person skilled in the art could have easily conceived of the idea of using replicon RNA where a genome of hepatitis C virus of genotype 2a is under control of a preferred regulatory sequence to transform host cells, whereby infectious hepatitis C virus particles and a cell strain capable of releasing such virus particles on a stable basis are obtained.

Therefore, a person skilled in the art could have introduced the genome of the said hepatitis C virus at a downstream point in relation to an IRES sequence and a marker gene, referring to the description in document 4; produced a vaccine using the obtained virus particles, referring to the

WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY

International application No.
PCT/JP2005/003232

Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement description in document 1; and produced a virus vector using the said replicon RNA, as required. It is not considered that the subject matters of the above claims would produce a particular effect. The subject matters of claims 1-5 do not appear to involve an inventive step in view of documents 2 and 4 cited in the ISR. It is considered that document 2 describes the genome of hepatitis C virus of genotype 2a. Document 4 mentions that, by introducing a genome of hepatitis C virus at a downstream point in relation to an IRES sequence and a marker gene, the said genome can be replicated in host cells. A person skilled in the art could have easily conceived of the idea of introducing the genome of hepatitis C virus of genotype 2a at a downstream point in relation to an IRES sequence and a marker gene to replicate the said genome in host cells. It is not considered that the subject matters of the above claims would produce a particular effect.

Docket No.: 1254-0321PUS1

(PATENT)

ADM/tmh

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

	Patent Application of: ji WAKITA et al.	
Application No.: 10/589,902		Confirmation No.: N/A
Filed	: August 17, 2006	Art Unit: N/A
For:	NUCLEIC ACID CONSTRUCT CONTAINING FULLLENGTH GENOME OF HUMAN HEPATITIS C VIRUS, RECOMBINANT FULLENGTH VIRUS GENOME- REPLICATING CELLS HAVING THE NUCLEIC ACID CONSTRUCT TRANSFERRED THEREINTO AND METHOD OF PRODUCING HEPATITIS C VIRUS PARTICLE	Examiner: Not Yet Assigned
<u>LE</u> T	TTER SUBMITTING ADDITIONAL DOCUME PHASE FOR A PCT APPL	
P.O. 1	PCT missioner for Patents Box 1450 andria, VA 22313-1450	
Sir:		
	Under the provisions of 27 C.F.R. 1.495, attach necessary for entering the national phase in contational application.	_
	Attached is a copy of the Notification of Missing I	Requirements (371 Formalities Letter).
\boxtimes	Attached is the Executed Declaration and Power o	f Attorney Original Photocopy.
	The specification attached to the executed Declar copy of the specification that was filed in the	

Birch, Stewart, Kolasch & Birch, LLP

Application No.: 10/589,902 August 17, 2006, including any amendments thereto (if applicable) filed on even date therewith. \boxtimes The undersigned hereby declares that "Attorney Docket No. 1254-0321PUS1" on page 1 of the attached Inventors' Declaration corresponds to Appl. No.10/589,902 filed ACID **CONSTRUCT** August 17, 2006 entitled "NUCLEIC CONTAINING FULLLENGTH GENOME OF HUMAN HEPATITIS C VIRUS, RECOMBINANT FULLLENGTH VIRUS GENOME-REPLICATING CELLS HAVING THE NUCLEIC CONSTRUCT TRANSFERRED THEREINTO AND METHOD ACID PRODUCING HEPATITIS C VIRUS PARTICLE." Attached is an English language translation of the above-identified application that was filed in a foreign language, which should be used as the copy for examination purposes. See the attached Translator's Verification; or The undersigned states that the English translation attached hereto is a true and correct translation of the application as originally filed in a foreign language. \Box Attached are 0 sheet(s) of drawings. Please substitute these replacement drawings for the corresponding - sheet(s) of drawings on file in the above-identified application. Attached are substitute claims commencing on a separate sheet in accordance with 37 C.F.R. § 1.75(h). П Attached is a substitute abstract commencing on a separate sheet in accordance with 37 C.F.R. § 1.72(b). Attached is a substitute specification that complies with 37 C.F.R. § 1.52. The substitute specification does not contain new matter.

Attached is a preliminary amendment.

2 ADM/tmh

Docket No.: 1254-0321PUS1

Application No.: 10/589,902 Docket No.: 1254-0321PUS1 Applicant claims small entity status under 37 C.F.R. § 1.27. \Box Attached is a Supplemental Application Data Sheet (ADS). П Submitted concurrently herewith under separate cover for recording is an Assignment. Attached is a Petition for Extension of Time. \boxtimes The Government Filing Surcharge for late filing of oath and/or declaration in the amount of \$130.00 in accordance with 37 C.F.R. §§ 1.494 and 1.492 was previously paid for concurrently with the filing of the application on August 17, 2006. Attached hereto is the fee transmittal listing the required fees. If necessary, the Commissioner is hereby authorized in this, concurrent, and future replies, to charge payment or credit any overpayment to our Deposit Account No. 02-2448 for any additional fees required under 37 C.F.R. § 1.16 or under § 1.17; particularly, extension of time fees. Respectfully submitted, Dated: December 13, 2006 Andrew D. Meikle Registration No.: 32,868 BIRCH, STEWART, KOLASCH & BIRCH, LLP 8110 Gatehouse Road Suite 100 East P.O. Box 747 Falls Church, Virginia 22040-0747 (703) 205-8000

Attorney for Applicant

Attachment(s)

3 ADM/tmh